

## 800 **Chapter 1. Introduction**

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802 **Convening Lead Authors:** A. R. Ravishankara, NOAA; Michael J. Kurylo, NASA;

803 Anne-Marie Schmoltnner, NSF

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805 Ozone (O<sub>3</sub>) is the triatomic form of oxygen. It is a key atmospheric trace gas that is  
806 present everywhere in the atmosphere and is most abundant in the stratosphere. The  
807 abundance of ozone in the stratosphere is largest in the region between roughly 15 and 35  
808 km, which is referred to as the stratospheric ozone layer. This stratospheric ozone layer  
809 (Box 1.1) plays many important roles in the Earth system:

- 810 • It protects the lower part of the atmosphere (the troposphere) and the Earth's  
811 surface from damaging, or "harsh" ultraviolet<sup>1</sup> (UV) radiation from the sun;
- 812 • It influences the chemical composition of the lower atmosphere by altering the  
813 amount and type (wavelength distribution) of solar radiation passing through it;
- 814 • It changes the temperature structure of the stratosphere and thus influences  
815 atmospheric transport and mixing; and
- 816 • It contributes ozone to the upper troposphere, where ozone is an important  
817 greenhouse gas.

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819 Because of many of the above contributions, ozone in the stratosphere and its changes  
820 also play a significant role in the Earth's climate system; changes in the ozone layer are  
821 influenced by climate change and also contribute to climate change. Appendix A of this

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<sup>1</sup> 'Harsh' UV radiation indicates the higher energy portion of the UV spectrum

822 report contains background information and the answers to some of the most frequently  
823 asked questions about the ozone layer (Fahey, 2007).

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825 The focus of this report is on key issues related to (a) the stratospheric ozone layer,  
826 including its changes in the past, its current abundances, and expected levels in the future;  
827 (b) emissions of ozone-depleting substances and their influences on the ozone layer and  
828 climate; and (c) the changes in the ground level UV radiation associated with  
829 stratospheric ozone changes.

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831 The chemical processes that lead to the formation of ozone as well as those that remove  
832 or destroy it, are distinctly different in the stratosphere from those in the troposphere  
833 (Box 1.2). The ever-present balance in the stratosphere between production, removal, and  
834 transport determines the abundance of ozone in any given part of the ozone layer. The  
835 majority of the removal processes in the stratosphere involve catalytic cycles in which  
836 ozone-destroying chemicals are reformed after destroying ozone. This catalytic capability  
837 is a key reason why very small amounts of ozone-destroying chemicals introduced into  
838 the atmosphere can vastly influence the ozone layer (Box 1.2).

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840 The potential for human-produced chemicals, such as chlorofluorocarbons (CFCs), to  
841 deplete the stratospheric ozone layer has received a great deal of attention since the early  
842 1970s. The depletion by chlorine released from CFCs in the stratosphere was expected to  
843 be catalytic in nature, in that small amounts of CFCs could destroy vast amounts of  
844 ozone. The ozone depletion was predicted to lead to changes in UV radiation at the

845 surface, with potentially major environmental consequences. The anticipated effects of  
846 increased UV radiation included increased incidence of skin cancer and cataracts in  
847 humans, detrimental effects on ecosystems including the aquatic system, and deleterious  
848 effects on materials, such as rubber and plastics. These potential effects were debated and  
849 the nations of the world agreed to protect the ozone layer through the 1985 Vienna  
850 Convention. Then the ozone hole in Antarctica was discovered in 1985. Investigation of  
851 the causes of this annually recurring polar springtime ozone depletion indicated that  
852 chlorofluorocarbons and other ozone-depleting chemicals were involved in additional  
853 catalytic ozone destruction pathways unique to the extremely cold polar stratosphere. It  
854 was also discovered that small particles containing water and nitric and/or sulfuric acid  
855 that are found in polar stratospheric clouds (PSCs) play a crucial role in these processes  
856 by converting chemically less reactive halogen-containing chemicals into more reactive  
857 chemicals, which are more effective in ozone depletion, and involved some catalytic  
858 cycles unique to this region.

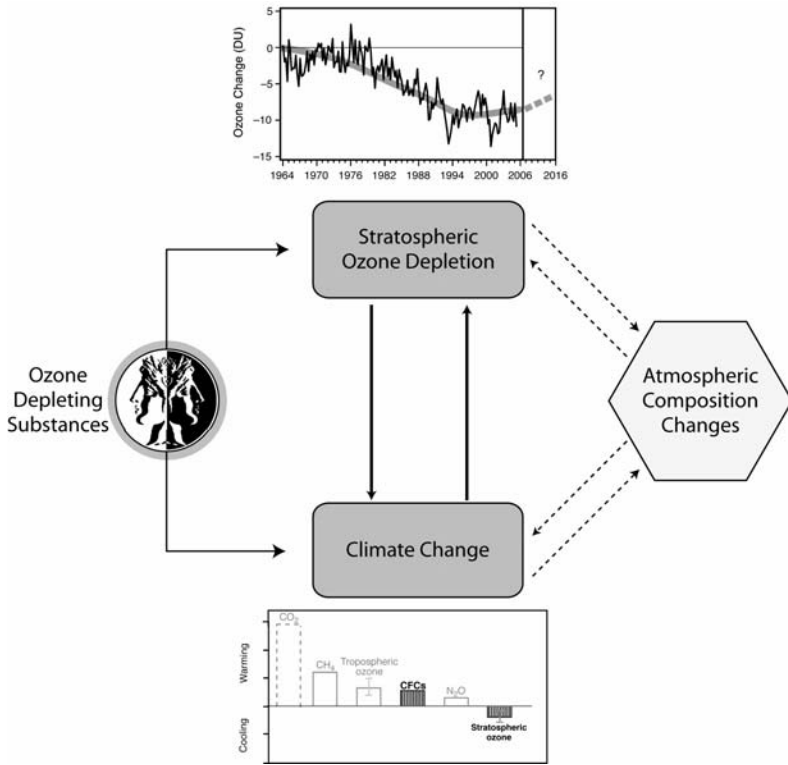
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860 The Montreal Protocol, a sequel to the Vienna convention, was agreed to in 1987 against  
861 the setting of the scientific knowledge at that date. First, the agreements of the Protocol  
862 were to reduce CFC emissions and to replace as much of the chlorofluorocarbons by the  
863 replacements that could be used in existing devices for most applications. A few  
864 applications utilized not-in-kind non-ozone-depleting chemicals. Many of the  
865 replacement chemicals still contained chlorine, but overall were less harmful to the  
866 stratospheric ozone layer than CFCs; many of these were hydrochlorofluorocarbons  
867 (HCFCs). Slowly, even the chlorine-containing substitutes were to be replaced by non-

868 chlorine or bromine containing replacements; many of these are hydrofluorocarbons  
869 (HFCs).

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871 During the course of the past three decades of ozone-layer research, it has become clearer  
872 that ozone-depleting substances, as well as many of the CFC-substitutes introduced to  
873 comply with the Montreal Protocol, are also potent greenhouse gases. Ozone depletion  
874 and climate change are distinct issues but are inextricably linked because ozone itself is a  
875 greenhouse gas and many of the ozone-depleting gases are potent greenhouse gases. To  
876 add to the complexity, changes in the major greenhouse gases such as carbon dioxide  
877 (CO<sub>2</sub>), methane (CH<sub>4</sub>), and nitrous oxide (N<sub>2</sub>O) also influence ozone depletion. Increases  
878 in CO<sub>2</sub> lead to a cooling of the stratosphere, which increases ozone in the upper  
879 stratosphere in non-polar regions, but decreases ozone in the polar lower stratosphere.  
880 The influence of CH<sub>4</sub> and N<sub>2</sub>O on the stratospheric ozone layer is dominated by their  
881 chemical interactions. Figure 1.1 captures this influence in a schematic form. An  
882 assessment of the climate effects of ozone-depleting substances has to consider both of  
883 their roles: as chemicals that deplete ozone, and as greenhouse gases that alter climate.  
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887 **Figure 1.1** The two faces of ozone-depleting substances: their roles as depleting agents of stratospheric  
 888 ozone, and as greenhouse gases that influence climate. The two roles are further interconnected because  
 889 ozone itself is a greenhouse gas and because climate change can lead to changes in the ozone layer. The  
 890 various connections between these two phenomena are shown. A plot of the changes in the observed global  
 891 ozone illustrates the stratospheric ozone depletion issue. The radiative forcing due to various greenhouse  
 892 gases, including ODSs, depicts the greenhouse gas issue and stratospheric ozone changes.  
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894 Since 1987, there have been many amendments and adjustments to the Montreal Protocol  
 895 to accelerate efforts to curtail the emissions of ozone-depleting substances (ODSs). These  
 896 actions have come about in response to our evolving knowledge of the ozone layer and its  
 897 changes, and have led to a reduction in the emissions and subsequently in the  
 898 atmospheric abundances of most ozone-depleting substances. Thus, the projected  
 899 extremely high atmospheric abundances of ODSs and the associated larger-scale  
 900 stratospheric ozone depletions were prevented from occurring. However, many key  
 901 questions remain:

- 902       • Are the emission controls working as anticipated, *i.e.*, are the atmospheric  
903       abundances of ozone-depleting substances declining as expected?
- 904       • Is the ozone layer recovering due to decreases in emissions of ODSs as predicted?
- 905       • Are the changes in UV occurring as expected with changes in ozone?
- 906       • What are the influences of other Earth system changes, *e.g.*, climate and  
907       atmospheric composition, on the ozone layer and its recovery from the ODS-  
908       induced depletion?
- 909       • What are the influences of ODSs, and their substitutes, on other aspects of the  
910       Earth system, especially climate?

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912   Because many ODSs have lifetimes of many years in the atmosphere, the depletion of  
913   stratospheric ozone is a global problem, and emissions of ODSs anywhere on the globe  
914   contribute to the ozone layer depletion. The extent of the ozone layer depletion for a  
915   given emission differs depending on the location (*e.g.*, latitude) and time (*e.g.*, season).  
916   Therefore, the observed ozone depletion in a given region will not be directly related to  
917   the emissions from that region. Yet, it is appropriate to ask: what is the contribution of  
918   one nation, or region, to the depletion of the global ozone layer? And, how do the ODSs  
919   influence stratospheric ozone, and hence UV, in a specific region or over a specific  
920   nation? Of course, it may not be feasible to answer these questions completely at the  
921   present time, given our current (and evolving) state of knowledge.

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923   This Synthesis and Assessment Product (SAP) of the Climate Change Science Program  
924   (CCSP), SAP 2.4, addresses key issues related to the stratospheric ozone layer, including

925 its changes in the past and its expected evolution in the future. Also, it takes account of  
926 the current abundances and emissions of ozone-depleting substances. Further, it  
927 synthesizes the best available information on the past and future levels of ultraviolet  
928 radiation at the Earth's surface. Lastly, it explores the interactions between climate  
929 change and stratospheric ozone changes as well as the ODS changes, and briefly recounts  
930 the influence of stratospheric ozone changes on climate change. All of these topics are  
931 carried out within the context of the United States of America to distill a regional  
932 assessment from current global assessments. More specifically, this document:

- 933 • Summarizes current quantitative information on sources (*i.e.*, emissions), sinks  
934 (*i.e.*, the removal pathways and their speed), and abundances of ozone-depleting  
935 substances and associated uncertainties; describes how the combined influence of  
936 chlorinated and brominated ODSs in the stratosphere can be quantified, and how  
937 all these are likely to change in the future.
- 938 • Discusses levels of ozone in various regions of the stratosphere, including the  
939 polar regions, paying special attention to the Antarctic ozone hole.
- 940 • Provides information on the past, current, and anticipated future levels of  
941 ultraviolet radiation.
- 942 • Provides an assessment of the impact of changes in both climate and atmospheric  
943 composition on the future of the ozone layer.
- 944 • Provides a brief assessment of the contribution of ozone-depleting substances on  
945 forcing of climate because these chemicals are also greenhouse gases.
- 946 • Describes how these findings relate to human activities, with a particular  
947 emphasis on the U.S. Special emphasis has been placed on quantifying the

948 contributions of the United States of America to the global amounts of ODSs.  
949 Further, given the influence that ODSs and substitute chemicals have on climate,  
950 the report attempts to calculate the contributions to the relief of climate change  
951 via reductions in the emissions of ODSs and switching over to more climate-  
952 friendly and ozone-friendly CFC substitutes.

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954 The primary sources of information for this report are the World Meteorological  
955 Organization (WMO) / United Nations Environment Programme (UNEP) 2006  
956 assessment on the ozone layer *Scientific Assessment of Ozone Depletion: 2006* (WMO,  
957 2007), and the 2005 Special Report of the Intergovernmental Panel on Climate Change  
958 (IPCC) on *Safeguarding the Ozone Layer and the Global Climate System – Issues*  
959 *Related to Hydrofluorocarbons and Perfluorocarbons* (IPCC/TEAP, 2005) and  
960 references therein. In addition, this report bases some findings on a few peer-reviewed  
961 publications of direct import to this issue that have become available since the  
962 finalization of the two international assessments. The report was initiated before the  
963 release of the IPCC Fourth Assessment Report (AR4). Therefore, this report does not rely  
964 on the IPCC AR4; however, some key pertinent issues from the IPCC report are used in a  
965 few instances where updated information was essential. They are noted as such in those  
966 chapters.

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971 **CHAPTER 1 REFERENCES**

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981 D. de Jager, T. Kestin, M. Manning, and L.A. Meyer (eds.)], 488 pp, Cambridge  
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983 **WMO** (World Meteorological Organization), 2007: *Scientific Assessment of Ozone*  
984 *Depletion: 2006*, Global Ozone Research and Monitoring Project—Report No. 50, 572  
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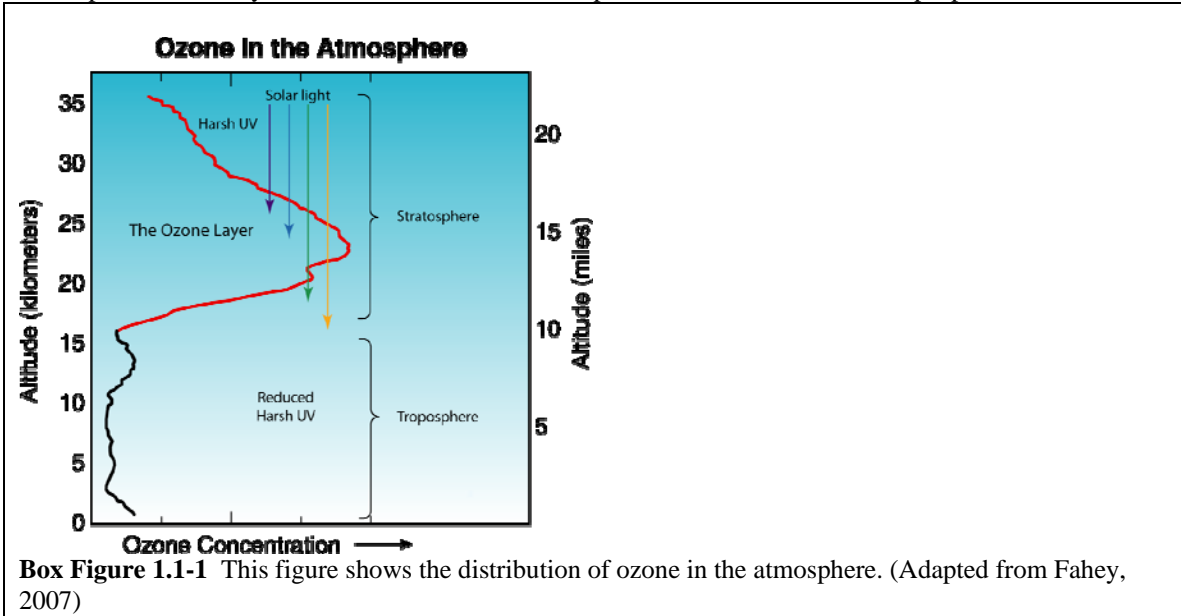
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**BOX 1.1: The Stratospheric Ozone Layer and its Role in the Atmosphere.**

About 90% of the atmospheric ozone resides in the stratosphere, in a region between roughly 15 and 35 km above the Earth’s surface, as indicated by the red line in Box Figure 1.1-1. This region is referred to as the stratospheric ozone layer. The remainder of the atmospheric ozone resides in the troposphere, the lower



**Box Figure 1.1-1** This figure shows the distribution of ozone in the atmosphere. (Adapted from Fahey, 2007)

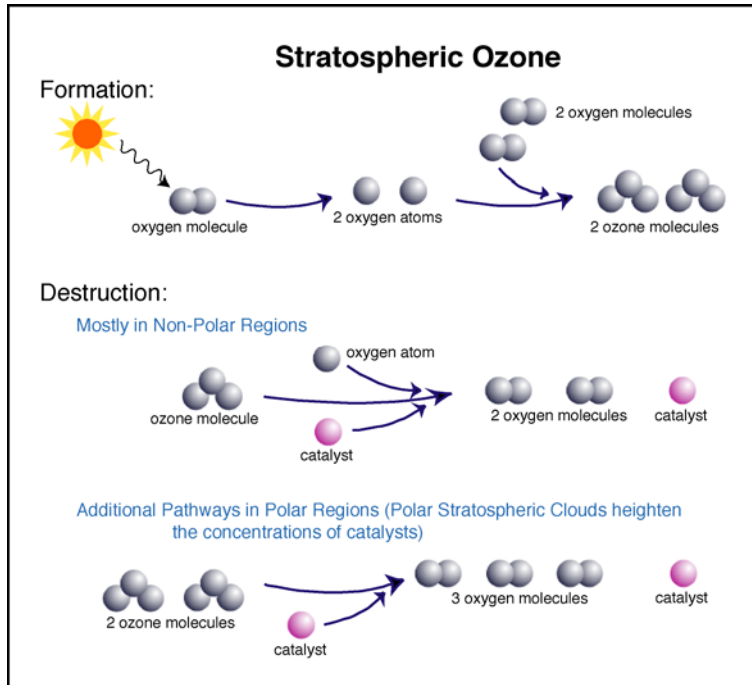
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layer of the atmosphere. Stratospheric ozone is formed and destroyed by chemical reactions, as shown in Box 1.2. Of particular note are the need for higher-energy UV radiation for the formation of ozone and the catalytic nature of the ozone removal processes. The ozone layer in turn shields the lower part of the atmosphere and the surface from damaging UV radiation because ozone itself absorbs UV radiation. Depletion of the ozone layer allows more UV- radiation (wavelength 280 to 315 nanometers) to reach the Earth’s surface. This radiation is harmful to humans and many other biological systems and causes damage to materials. The ozone in the lower atmosphere, the troposphere, is formed by methods different from those in the stratosphere, as shown in Box 1.2. Further, the contribution of this lower atmospheric ozone to the total in the atmosphere is small, of the order of a few percent in the southern hemisphere to about 10% in the northern hemisphere. The ozone in the lower atmosphere is harmful because, in direct contact, ozone is toxic to biological systems and can deteriorate many materials. It can cause respiratory and other health problems for humans. In addition, ozone and its changes in both the stratosphere and the lower atmosphere are important greenhouse gases and thus their changes influence climate. See Appendix A of this Synthesis and Assessment Product for further background information about ozone.

\*\*END BOX 1.1 \*\*\*\*\*

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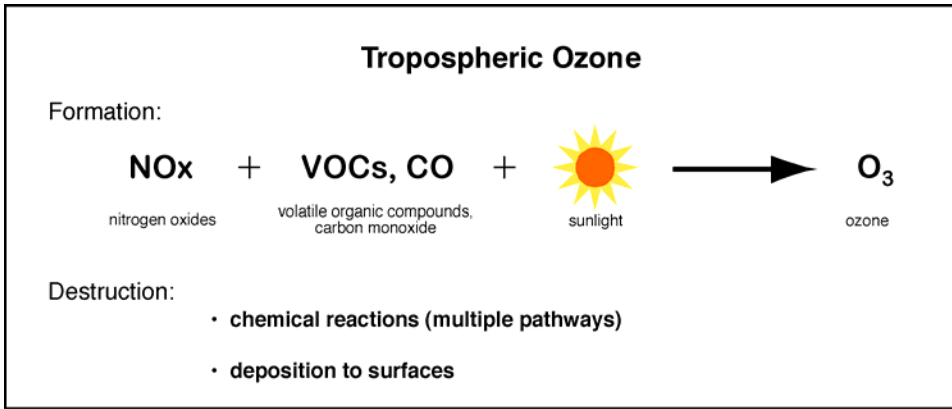
**BOX 1.2: A Simplified Representation of the Production and Removal of Ozone in the Atmosphere - the Processes that Determine the Abundance of Ozone.**



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**Box Figure 1.2-1** Highly simplified representation of the chemical processes that lead to the production and removal of ozone in the stratosphere. See Chapter 3 and Appendix A for further details.

Oxygen molecules ( $O_2$ ) are broken apart by the harsh UV radiation in the stratosphere to produce atomic oxygen, which reacts further with oxygen molecules to make ozone ( $O_3$ ). The ozone in the stratosphere is removed predominantly via catalytic chemical reactions that regenerate the catalysts. The catalysts include atoms and radicals produced in the stratosphere from the breakdown of various chemicals emitted at the Earth's surface. They include naturally occurring chemicals such as nitrogen oxides and hydrogen oxides, as well as human-emitted chemicals containing chlorine and bromine atoms, such as chlorofluorocarbons (CFCs) and bromine-containing halons that are used as fire extinguishants. These human-emitted species, referred to as ozone-depleting substances (ODSs), are of concern for the depletion of the ozone layer. The destruction pathway marked "non-polar regions" in Box Figure 1.2-1 is predominant outside of the springtime polar regions, while the pathway marked "polar regions" is dominant in the springtime polar ozone depletion including the Antarctic ozone hole. Because of the nature of these chemical processes, as discussed above, a very small amount of the catalyst (for example, chlorine atoms from CFCs) can destroy a large amount of stratospheric ozone. In addition to these chemical processes, transport of ozone (redistribution) is key to determining the abundance of ozone in a given location.



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**Box Figure 1.2-2** Schematic representations of the chemical processes that lead to the production and removal of ozone in the troposphere.

In contrast to the stratosphere, in the troposphere ozone is made using near UV and visible radiation (*i.e.*, longer wavelength) because the higher energy, harsh UV (shorter wavelength) is screened out by the stratospheric ozone layer. This tropospheric ozone production process requires nitrogen oxides, mostly from combustion, and volatile organic compounds. Unlike stratospheric ozone, tropospheric ozone is removed not only by chemical reactions but also by other processes including contact with the surface. The transport of ozone from the stratosphere to the troposphere is important as an ozone source in certain regions.

\*\*\*\*\*END BOX 1.2 \*\*\*\*\*